## **Amendments to the Claims**

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The following listing of claims replaces all previous listings or versions thereof:

- 1. (Currently amended) Method Method for detecting endotoxin, comprising the steps:
  - a) incubation of incubating a sample with a bacteriophage tail protein, and
  - b) detection of detecting endotox in bonded to bacteriophage tail proteins.
- 2. (Currently amended) Method The method according to claim 1, if necessary further comprising furthermore after step a) and prior to step b) the additional step of:
  - a') separation of separating the bacteriophage tail protein-endotoxin complexes from the sample.
- 3. (Currently amended) Method The method according to one of the claims claim 1 to 3, the detection being implemented by means of wherein detection comprises spectroscopic methods.
- 4. (Currently amended) Method for removing endotoxin from a sample, comprising the steps:
- a) incubation of incubating a sample with or bringing a sample in contact with bacteriophage tail proteins which are immobilised on a permanent carrier, non specifically or directed,
- b) separation of separating the bacteriophage tail protein-endotoxin complex from the sample.

- 5. (Currently amended) Method The method according to claim 4, the wherein steps a) and b) being are implemented in a chromatography column throughflow method.
- 6. (Currently amended) Method The method according to claim 4, wherein the permanent carrier being comprises filtration media, glass particles, magnetic particles, centrifugation materials, sedimentation materials or filling materials for chromatography columns.
- 7. (Currently amended) <u>Method The method</u> according to claim 4-to-6, the bacteriophage tail proteins being immobilised on the permanent carrier via coupling groups.
- 8. (Currently amended) Method The method according to claim 7, the coupling group being a lectin, receptor or anticalin.
- 9. (Currently amended) <u>Method The method</u> according to claim 7, <u>wherein</u> the coupling group <u>being-acomprises</u> streptavidin or avidin and the bacteriophage tail proteins <u>beingis</u> coupled with biotin or a Strep-tag.
- 10. (Currently amended) Method The method according to claim 4-to-6, the bacteriophage tail proteins being immobilised on the permanent carrier covalently via chemical bonds.
- 11. (Currently amended) Method The method according to one of the preceding elaimsclaim 1, wherein the bacteriophage tail protein having comprises a Strep-tag or a Histag.

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- 12. (Currently amended) Method The method according to claim 11, wherein the tag having comprises an amino acid sequence according to SEQ ID NO. 5, 6 or 7.
- 13. (Currently amended) Method The method according claim 11-or 12, wherein the p12 protein of the phage T4 being is used as bacteriophage tail protein.
- 14. (Currently amended) Method The method according to one of the preceding elaimsclaim 1, wherein the Ca<sup>2+</sup> concentration inof the incubation being comprises 0.1  $\mu$ M to 10 mM and the Mg<sup>2+</sup> concentration being comprises 0.1  $\mu$ M to 10 mM.
- 15. (Currently amended) Method The method according to one of the elaims 1 to 3 claim 1, marked endotoxin being displaced from the bond with a bacteriophage tail protein and the marked endotoxin being subsequently detected.
- 16. (New) The method according to claim 1, wherein the bacteriophage tail protein comprises a Strep-tag or a His-tag.
- 17. (New) The method according to claim 16, wherein the tag comprises an amino acid sequence according to SEQ ID NO. 5, 6 or 7.
- 18. (New) The method according claim 16, wherein the p12 protein of the phage T4 being used as bacteriophage tail protein.

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